AMENDMENTS TO THE CLAIMS:

This listing of claims will replace all prior versions, and listings, of claims in the application:

- 1-5 (Cancelled).
- 6. (Previously Presented) A bioreductive conjugate of the formula II:

$$R^{1}$$
 R^{2}
 R^{2}
 R^{2}
 R^{2}
 R^{3}
 R^{2}
 R^{3}
 R^{2}
 R^{3}
 R^{3}
 R^{3}
 R^{3}
 R^{3}
 R^{3}
 R^{3}
 R^{3}

(wherein

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R¹ and R² independently represent hydrogen or halogen atoms, or a group R, OR, SR, NHR,

NR₂, CO₂R or CONHR;

or, alternatively, R1 and R2 together with the intervening ring carbon atoms form a 5-7

membered carbocyclic or heterocyclic ring itself optionally substituted by one or more halogen

atoms, or by one or more groups selected from R, OR, SR, NHR, NR2, CO2R and CONHR;

Z represents an alkyl, alkenyl, aryl or aralkyl group optionally carrying at least one OH, SH, NH₂

or NHR⁷ group in which R⁷ is an alkyl group or Z represents a group of the formula -XH where

X represents an oxygen or a sulphur atom, or a group of formula NY in which Y represents a

hydrogen atom or an alkyl group;

R³, R⁴, R⁵ and R⁶ independently represent hydrogen atoms or an alkyl or alkenyl group;

each group R independently represents a hydrogen atom, an alkyl or alkenyl group;

E represents the residue of a therapeutic agent to be delivered, optionally attached via a linking

group L which is an ester, phosphate ester, ether, amine, thiol or thiol ester group or any

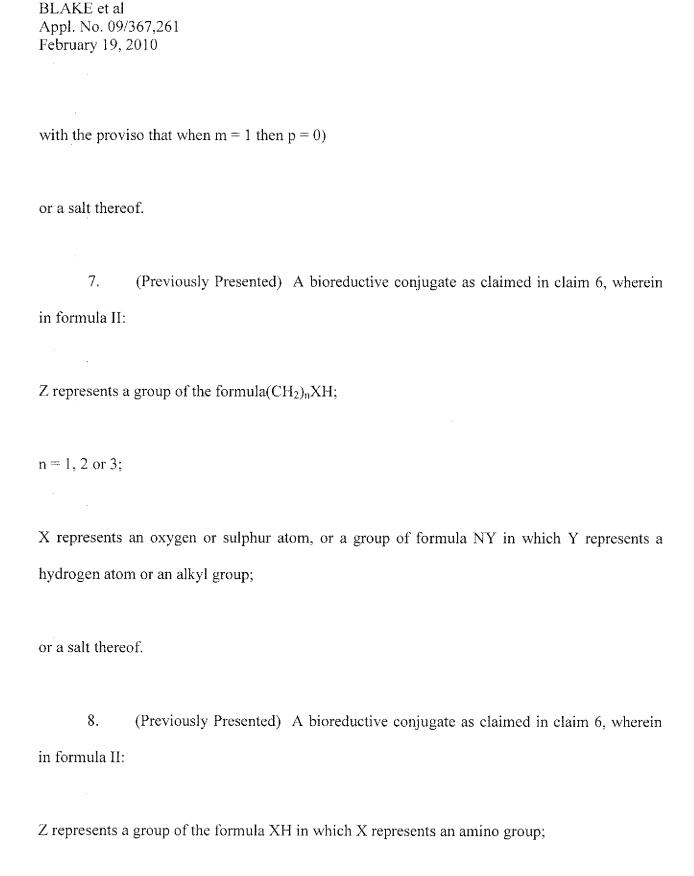
combination thereof;

m = 0, 1, 2 or 3; and

p = 0 or 2;

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 R^1 and R^2 each represent alkoxy groups or, together with the intervening ring carbon atoms, R^1 and R^2 form a benzene ring;

R³, R⁴, R⁵ and R⁶ each represent hydrogen atoms; and

$$n = m = 1$$
 and $p = 0$;

or a salt thereof.

9. (Withdrawn) A bioreductive conjugate of formula III:

(wherein

P and Q together with the intervening ring carbon atoms form a quinone or indoloquinone ring, an N-oxide or diazoaromatic compound, itself optionally substituted by one or more halogen atoms, or by one or more groups selected from R, OR, SR, NHR, NR₂, CO₂R and CONHR;

 R^1 represents a hydrogen or halogen atom, or a group R, OR, SR, NHR, NR_2 , CO_2R or CONHR;

R³, R⁴ and R⁵ independently represent hydrogen atoms or an alkyl or alkenyl group;

each group R independently represents a hydrogen atom, an alkyl or alkenyl group; and

E represents the residue of a therapeutic agent to be delivered, optionally attached via a linking group L which is an ester, phosphate ester, ether, amine, thiol or thiol ester group or any combination thereof;)

or a salt thereof.

10. (Withdrawn) A bioreductive conjugate as claimed in claim 9, wherein in formula III:

P and Q together with the intervening ring carbon atoms form a quinone or indoloquinone ring; and

R¹, R³, R⁴ and R⁵ each represent hydrogen atoms or methyl groups;

or a salt thereof.

11. (Withdrawn) A bioreductive conjugate of formula IV:

$$\begin{array}{c|c}
 & E \\
 & R5 \\
 & R7 \\
 & Z
\end{array} (IV)$$

(wherein

S and T together with the intervening ring carbon atoms form a quinone or indoloquinone ring, an N-oxide compound, itself optionally substituted by one or more halogen atoms, or by one or more groups selected from R, OR, SR, NHR, NR₂, CO₂R and CONHR;

Z represents an alkyl, alkenyl, aryl or aralkyl group optionally carrying at least one OH, SH, NH_2 or NHR^6 group in which R^6 is an alkyl group;

R⁷ represents an alkyl group;

R³, R⁴ and R⁵ independently represent hydrogen atoms or an alkyl or alkenyl group;

each group R independently represents a hydrogen atom, an alkyl or alkenyl group;

q = 0, 1, 2 or 3; and

E represents the residue of a therapeutic agent to be delivered, optionally attached via a linking group L which is an ester, phosphate ester, ether, amine, thiol or thiol ester group or any combination thereof;)

or a salt thereof.

12. (Withdrawn) A bioreductive conjugate as claimed in claim 11, wherein in formula IV:

S and T together with the intervening ring carbon atoms form a quinone or N-oxide compound;

R³, R⁴ and R⁵ each represent hydrogen atoms;

R⁷ is methyl;

Z represents a group of formula (CH₂)_nXH wherein X represents an oxygen or sulphur atom, or X represents a group of formula NY in which Y represents a hydrogen atom or an alkyl group; and

$$q = 0 \text{ or } 1,$$

or a salt thereof.

13-16 (Cancelled).

17. (Previously Presented) A bioreductive conjugate as claimed in claim 6 wherein said linker group L if present is a group of the formula:

$$O-CO-(CH_2)_n-CO-X-$$

or

(wherein n is an integer from 1 to 3;

X represents a sulphur or oxygen atom; and

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R⁸ and R⁹ each independently represent F or Cl).

18 and 19 (Cancelled).

20. (Previously Presented) A pharmaceutical composition comprising a bioreductive

conjugate as claimed in claim 6, or a pharmaceutically acceptable salt thereof, together with at

least one pharmaceutical carrier or excipient.

21. (Previously Presented) A bioreductive conjugate as claimed in claim 6 for use in

a method of targeting a therapeutic agent to a site of hypoxia and/or ischemia within the human

or non-human animal body.

22. (Currently Amended) A method of treating bioreductive conjugate as claimed in

claim 6 for use in the treatment of rheumatoid arthritis or other arthritic conditions, diabetes,

atherosclerosis, stroke, sepsis, Alzheimer's disease and other neurological disorders, cancer,

kidney disease, digestive diseases, liver disease, chronic periodontitis or ischemia following

tissue transplantation comprising administering to a patient in need thereof an amount of the

bioreductive conjugate as claimed in claim 6 sufficient to effect said treatment.

23. (Cancelled).

24. (Currently Amended) A method of treating Use as claimed in claim 22 for the

treatment of rheumatoid arthritis or other arthritic conditions, diabetes, atherosclerosis, stroke,

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sepsis, Alzheimer's disease and other neurological disorders, cancer, kidney disease, digestive diseases, liver disease, chronic periodontitis or ischemia following tissue transplantation comprising administering to a patient in need thereof an amount of the bioreductive conjugate as claimed in claim 9 sufficient to effect said treatment.

- 25. (Previously Presented) A method of targeting hypoxic and/or ischemic tissues in the human or non-human animal body, said method comprising administering to said body a bioreductive conjugate as claimed in claim 6.
- 26. (Withdrawn) A bioreductive conjugate as claimed in claim 9 wherein said linker group L if present is a group of the formula:

or

(wherein n is an integer from 1 to 3;

X represents a sulphur or oxygen atom; and

R⁸ and R⁹ each independently represent F or Cl).

27. (Withdrawn) A bioreductive conjugate as claimed in claim 11 wherein said linker group L if present is a group of the formula:

or

(wherein n is an integer from 1 to 3;

X represents a sulphur or oxygen atom; and

R⁸ and R⁹ each independently represent F or Cl).

28. (New) A method of treating rheumatoid arthritis or other arthritic conditions, diabetes, atherosclerosis, stroke, sepsis, Alzheimer's disease and other neurological disorders, cancer, kidney disease, digestive diseases, liver disease, chronic periodontitis or ischemia following tissue transplantation comprising administering to a patient in need thereof an amount of the bioreductive conjugate as claimed in claim 11 sufficient to effect said treatment.